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(54) **SYNTHESIS OF JANUS NANOMATERIALS**

FOREIGN PATENT DOCUMENTS

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(52) **U.S. Cl.**  
CPC ..... **C01B 33/12** (2013.01)

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None  
See application file for complete search history.

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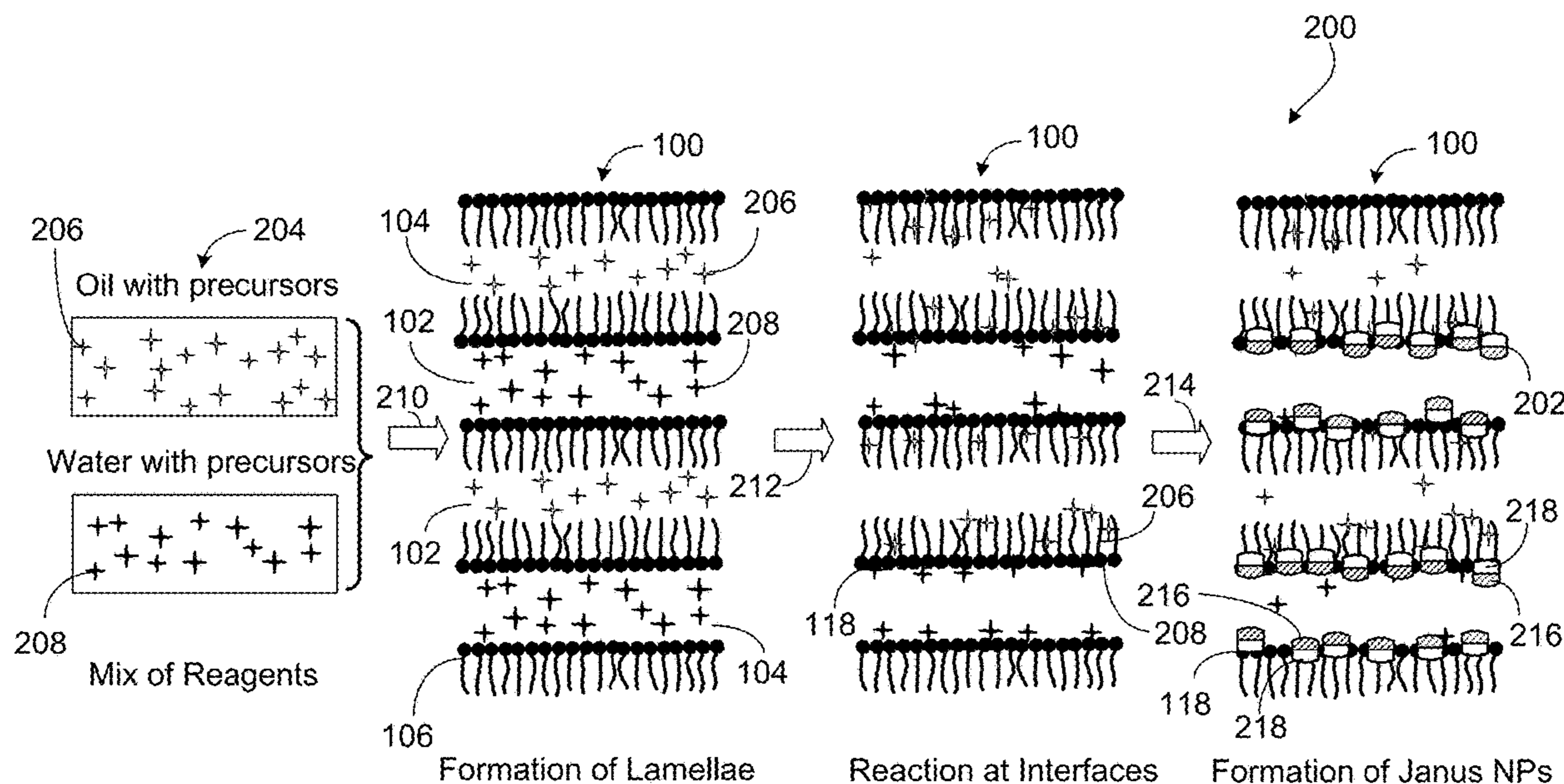
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(57) **ABSTRACT**

Synthesizing Janus nanoparticles including forming a lamellar phase having water layers, organic layers, and a surfactant, and reacting chemical precursors in the lamellar phase to form the Janus nanoparticles at interfaces of the water layers with the organic layers.

**8 Claims, 7 Drawing Sheets**



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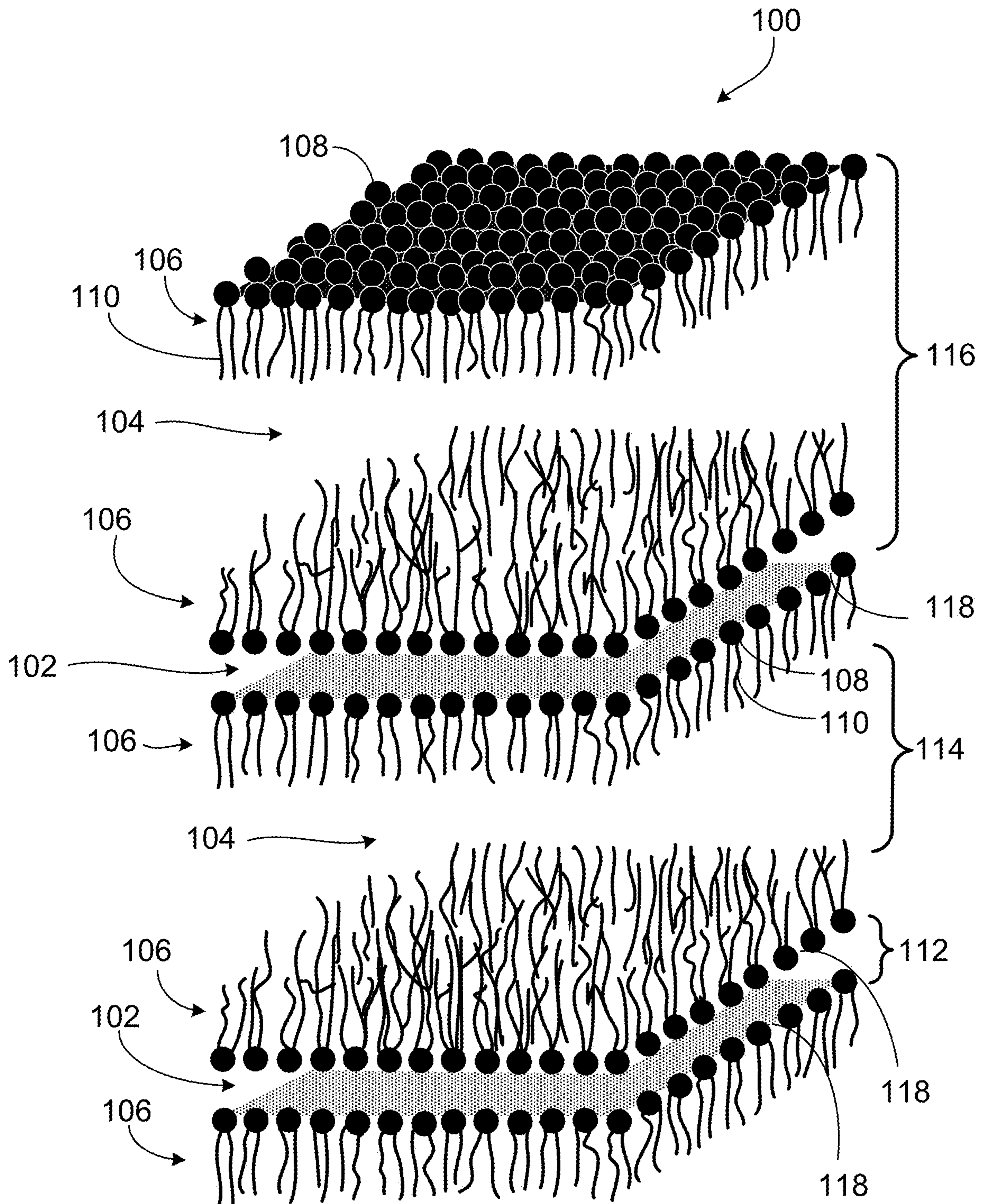


FIG. 1

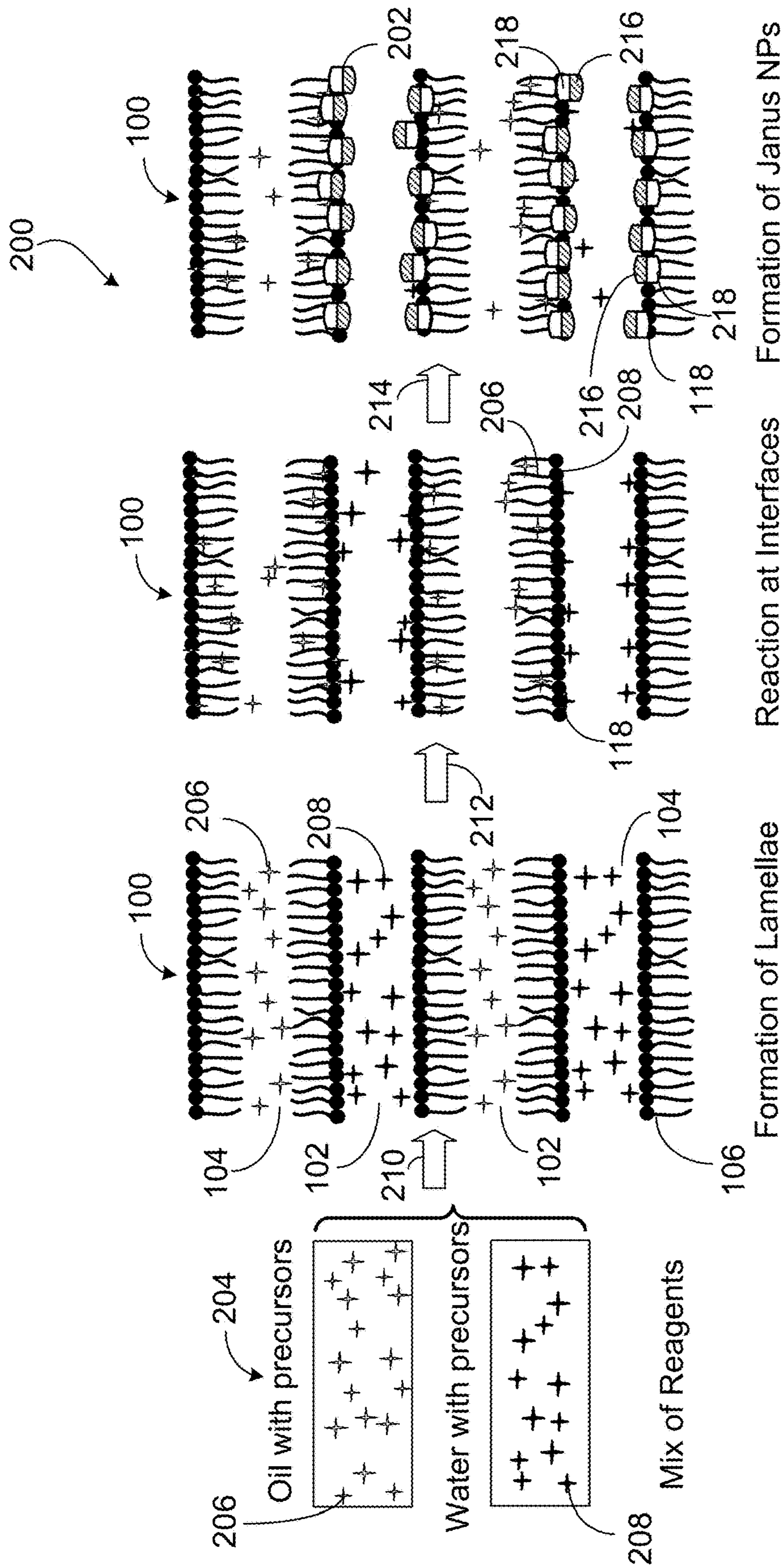
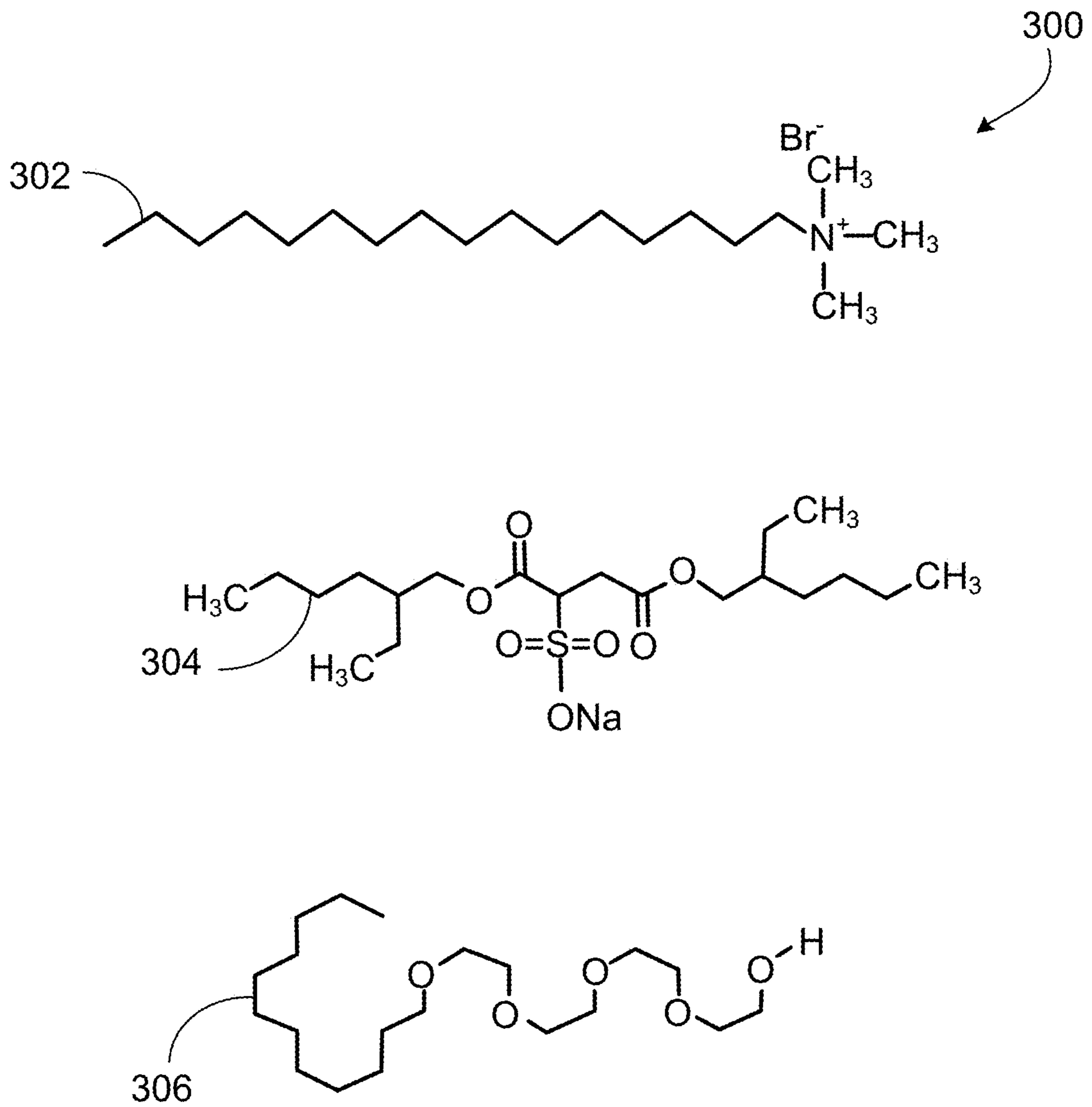


FIG. 2



**FIG. 3**

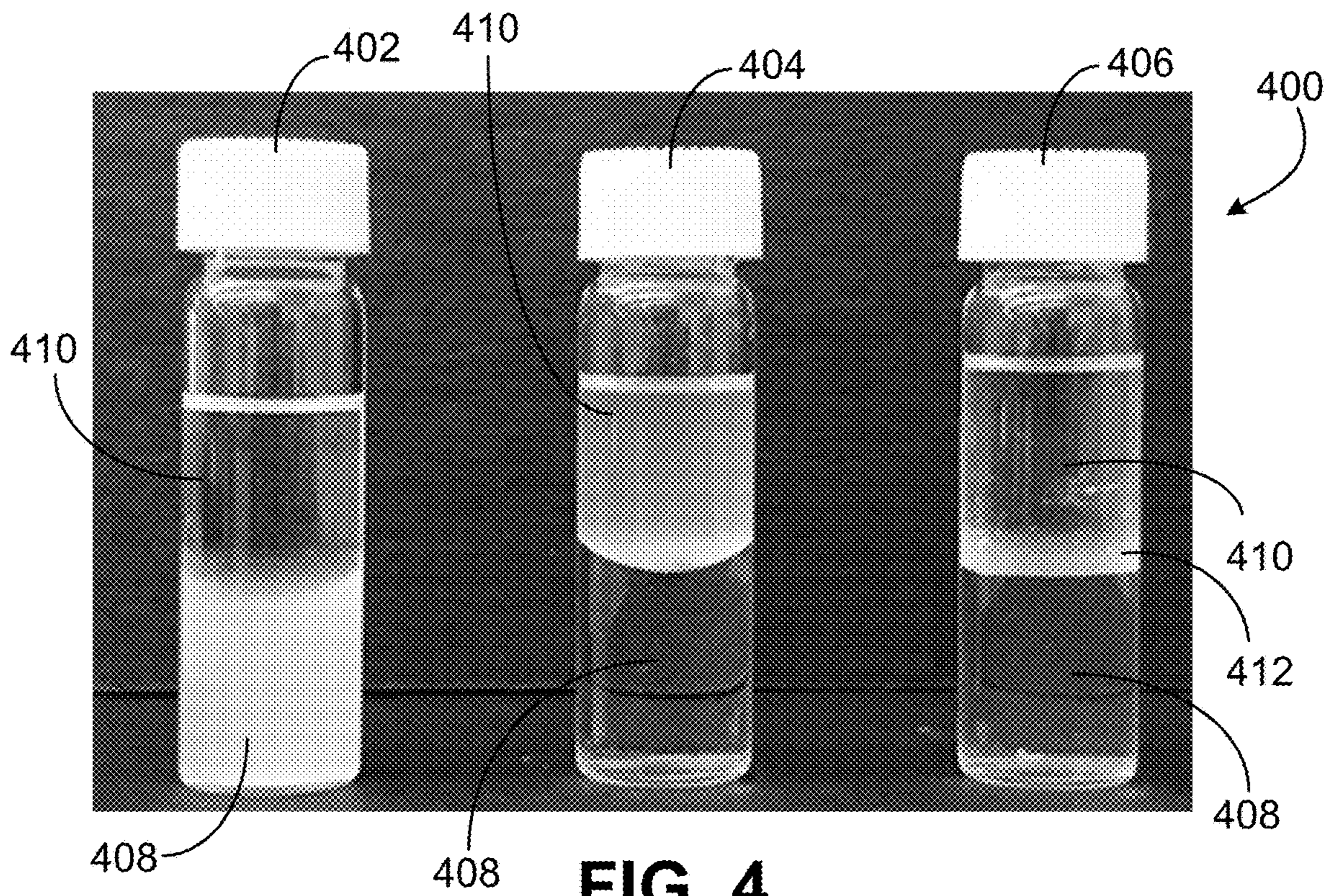


FIG. 4

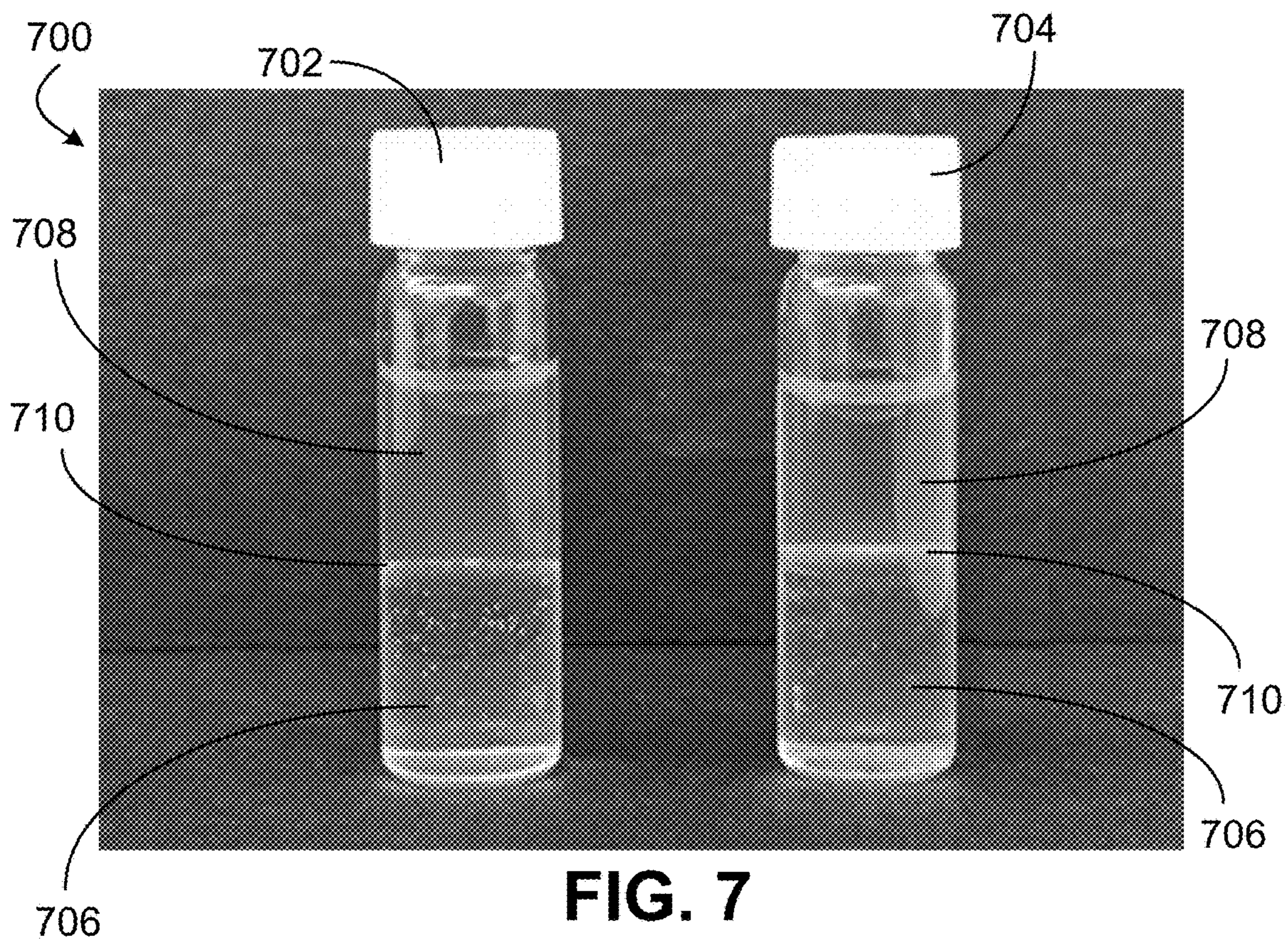


FIG. 7



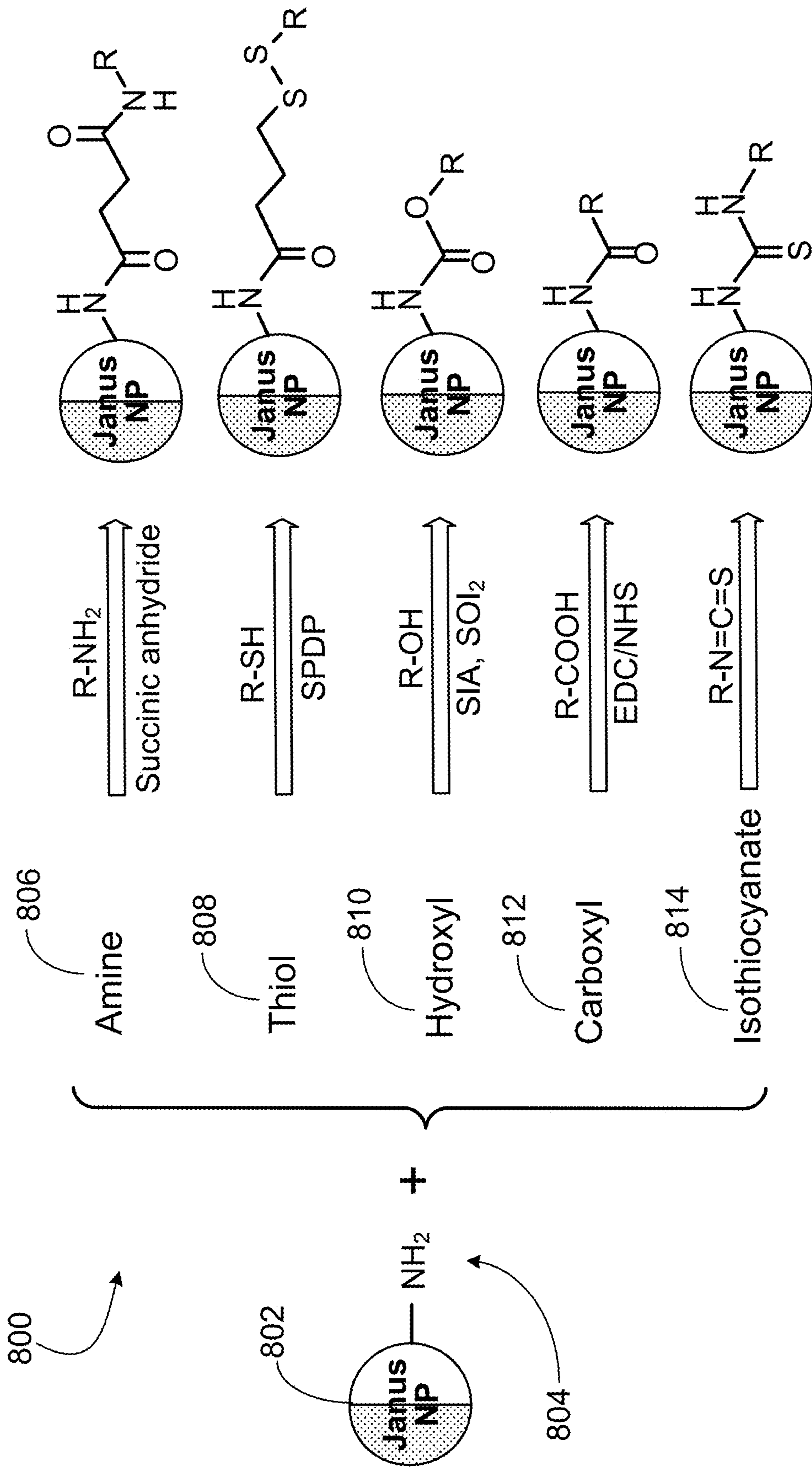


FIG. 8



900

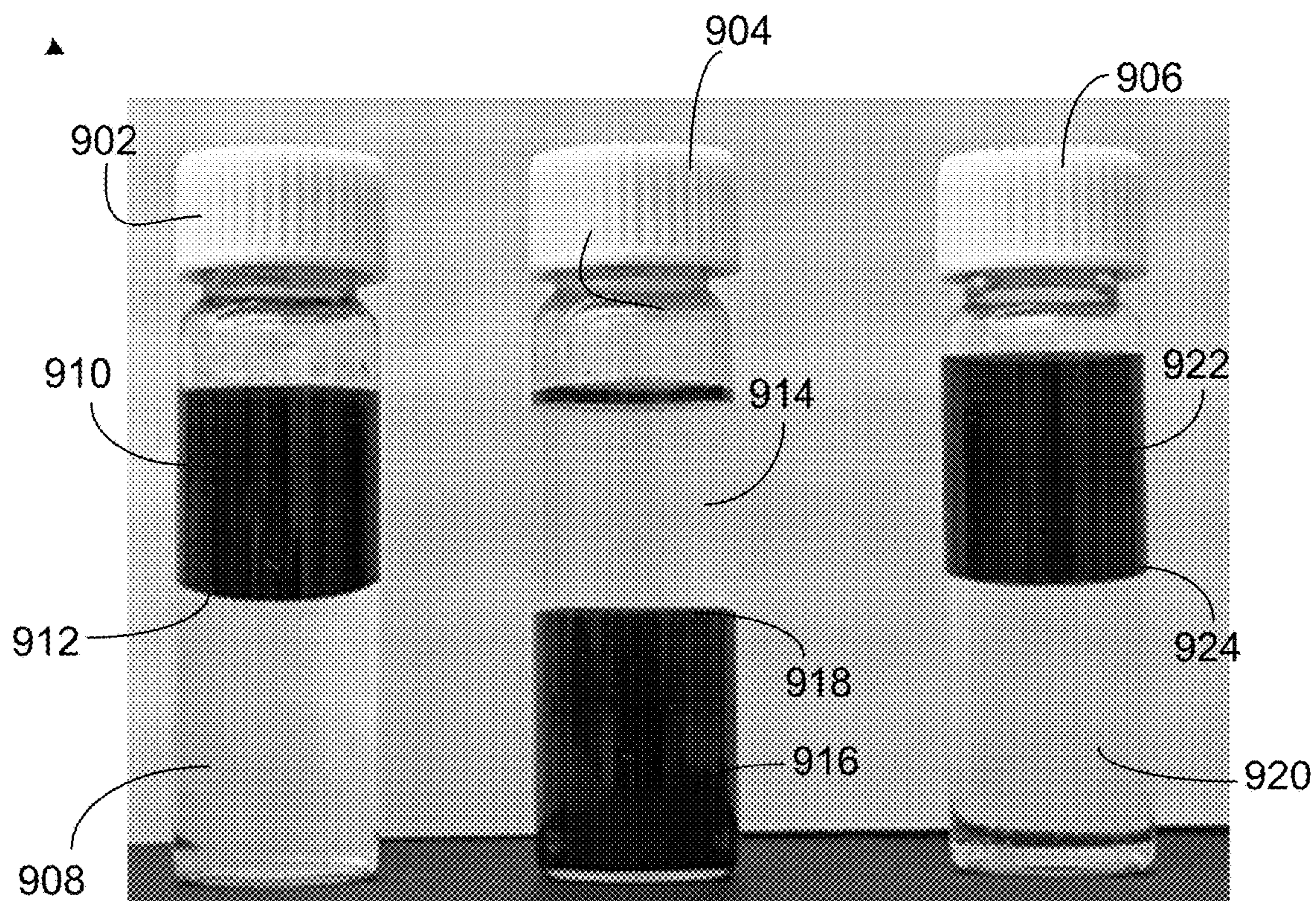


FIG. 9

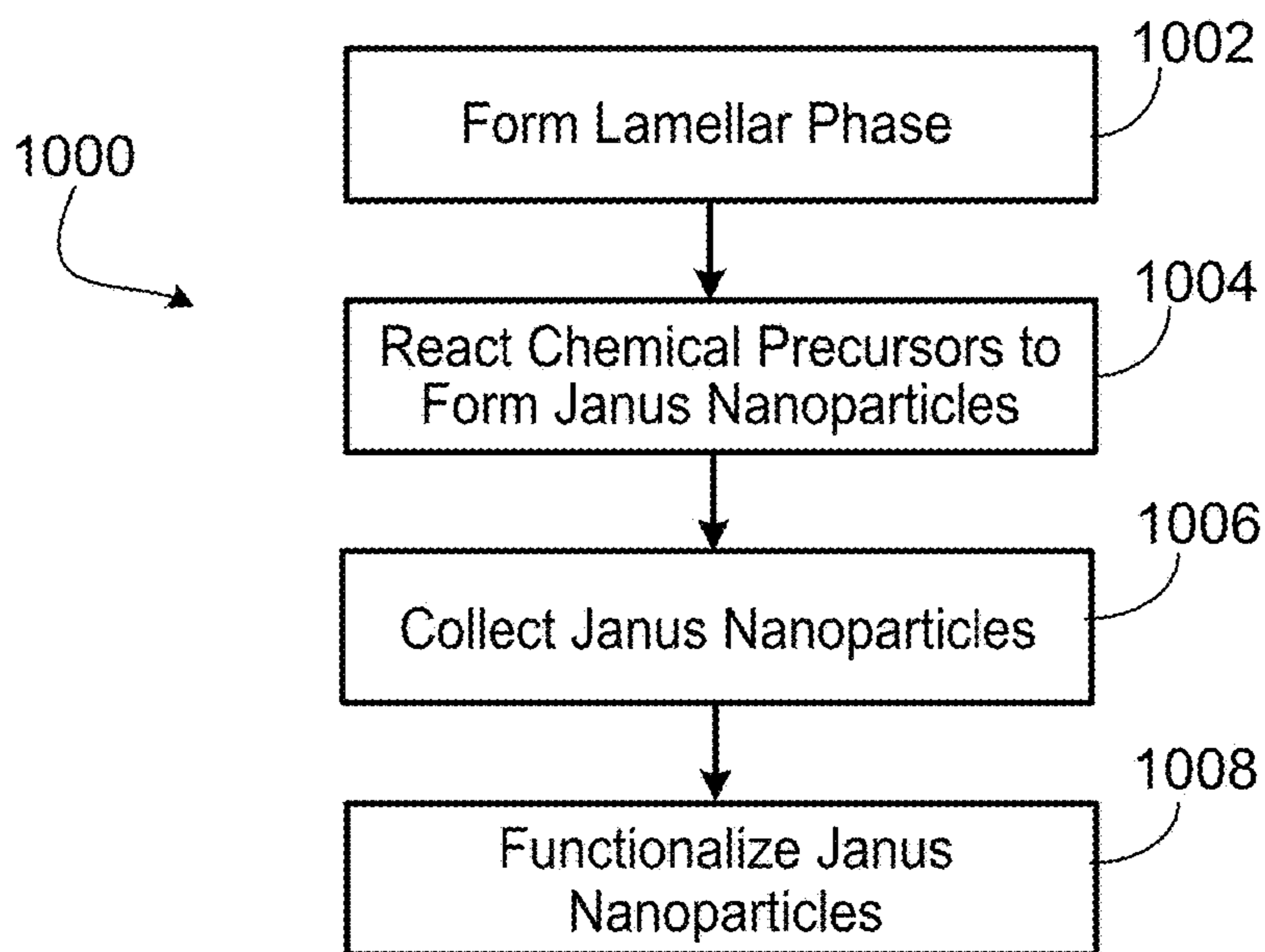


FIG. 10

## SYNTHESIS OF JANUS NANOMATERIALS

## TECHNICAL FIELD

This disclosure relates to Janus nanomaterials, such as Janus nanoparticles.

## BACKGROUND

Janus particles are types of nanoparticles with at least two surfaces having different respective physical properties. This surface arrangement of Janus nanoparticles allows two different types of chemistry to occur on the same particle. This surface configuration gives properties related to the asymmetric structure or asymmetric functionalization of the particles. The synthesis of Janus nanoparticles generally relies on selectively creating each side of a nanometer-sized particle with different chemical properties. Example syntheses of Janus nanoparticles include masking, self-assembly, and phase separation. Janus nanoparticles have generally been produced in the laboratory in small quantities, such as less than 100 milligrams (mg) in a day. Such syntheses typically employ multiple-step reactions.

## SUMMARY

An aspect relates to a method of synthesizing Janus nanoparticles. The method includes forming a lamellar phase having water layers, organic layers, and a surfactant. The method includes reacting chemical precursors in the lamellar phase to form Janus nanoparticles at interfaces of the water layers with the organic layers.

Another aspect relates to a method of synthesizing Janus nanoparticles. The method includes forming a lamellar phase having a surfactant, water layers, and organic layers. The method includes incorporating a first chemical reagent in the water layers and incorporating a second chemical reagent in the organic layers. The method includes allowing the first chemical reagent to react with the second chemical reagent at interfaces of the water layers with the organic layers to form Janus nanoparticles at the interfaces.

Yet another aspect relates to a method of synthesizing Janus nanomaterial. The method includes forming a lamellar phase having a surfactant, water layers, organic layers, and chemical precursors. The method includes allowing the chemical precursors to react to form Janus nanoparticles at interfaces between the organic layers and the water layers.

Yet another aspect relates to a lamellar phase having surfactant, water layers with a water-soluble organometallic compound, and oil layers with an oil-soluble organometallic compound. The lamellar phase has Janus metal-oxide nanoparticles formed from reaction of the water-soluble organometallic compound with the oil-soluble organometallic compound at the interfaces of the water layers with the oil layers. The Janus metal-oxide nanoparticles are disposed at the interfaces.

The details of one or more implementations are set forth in the accompanying drawings and the description later. Other features and advantages will be apparent from the description and drawings, and from the claims.

## BRIEF DESCRIPTION OF DRAWINGS

FIG. 1 is a diagram of a liquid-crystal lamellar phase having a water phase (water layers) and an organic phase (organic layers).

FIG. 2 is a sequence diagram of a lamellar phase in the synthesis of Janus nanoparticles.

FIG. 3 is a diagram giving exemplary surfactants utilized to form the lamellar phase.

FIG. 4 is an image of three vials each having different respective nanoparticles in hexane and water.

FIG. 5 is scanning electron microscope (SEM) images of Janus silicon-dioxide (SiO<sub>2</sub>) nanoparticles.

FIG. 6 is a diagram of a reaction giving formation of an isothiocyanate-amine complex.

FIG. 7 is an image of two vials each having Janus SiO<sub>2</sub> nanoparticles in a bi-phase mixture of chloroform and water.

FIG. 8 is a diagram depicting exemplary reactions that may further functionalize the Janus nanoparticles.

FIG. 9 is an image of three vials each having water and crude oil.

FIG. 10 is a block flow diagram of a method of synthesizing Janus nanoparticles.

Like reference numbers and designations in the various drawings indicate like elements.

## DETAILED DESCRIPTION

In some implementations, a Janus nanoparticle may be formed by dividing a nanoparticle into two distinct parts each of a different respective material or bearing different functional groups. For example, a Janus nanoparticle may have one-half of its surface composed of hydrophilic groups and the other half hydrophobic groups. This may impart properties related to the asymmetric structure or functionalization of the Janus nanoparticle. The synthesis of Janus nanoparticles may rely on selectively forming each side of a nanometer-sized particle with different chemical properties.

Janus particles in nanoscale (or submicron scale) may be unique among nanoscale materials because Janus particles provide asymmetry. The Janus particles are asymmetric. The Janus particles via their asymmetry can impart different chemical or physical properties and directionality within a single particle. The broken symmetry may provide for targeting self-assembled materials and realizing properties unavailable with homogeneous particles or symmetric patchy particles. Janus nanoparticles may have distinct properties.

Janus nanoparticles have generally not been manufactured in industrial-scale quantities. Janus nanoparticles have been synthesized at the laboratory research scale. Embodiments of the present techniques synthesize Janus nanomaterials at larger than research scale, such as greater than one kilogram (kg) in a day. Therefore, Janus nanomaterials may be produced at commercial scale for supply in applications across several industries. One example of an application is in the oil industry in which Janus nanoparticles, such as Janus ceramic nanoparticles or Janus silica nanoparticles, are utilized in nanofluid flooding in enhanced oil recovery (EOR). Aspects of the present techniques may provide for mass production of Janus nanomaterials or Janus nanoparticle materials for industrial application.

The synthesis or a portion of the synthesis of the Janus nanoparticles may be a one-pot synthesis. In chemistry, a one-pot synthesis may be that chemical reactions occur in one reactor. This may be desirable in avoiding separation and purification of intermediate chemical compounds and thus save time and resources.

The present disclosure relates to the high-throughput bulk synthesis (for example, greater than 1 kilogram per day) of Janus nanomaterials. Some aspects are directed to the high-

throughput bulk synthesis of Janus ceramic nanoparticles or Janus silica nanoparticles. A one-pot synthesis or one-pot mass synthesis of the Janus nanoparticles (for example, ceramic or silica) may be implemented.

The Janus ceramic particles synthesized may be Janus metal-oxide nanoparticles, such as Janus silicon-dioxide ( $\text{SiO}_2$ ) nanoparticles, Janus titanium-dioxide ( $\text{TiO}_2$ ) nanoparticles, Janus zirconium-dioxide ( $\text{ZrO}_2$ ) nanoparticles, Janus aluminum(III)-oxide ( $\text{Al}_2\text{O}_3$ ) nanoparticles, Janus iron(III)-oxide ( $\text{Fe}_2\text{O}_3$ ) nanoparticles, Janus zinc-oxide ( $\text{ZnO}$ ) nanoparticles, Janus tin-dioxide ( $\text{SnO}_2$ ) nanoparticles, or Janus gallium(III)-oxide ( $\text{Ga}_2\text{O}_3$ ) nanoparticles. The Janus metal-oxide nanoparticles can be Janus rare-earth metal-oxide nanoparticles, such as Janus cerium(IV)-oxide ( $\text{CeO}_2$ ) nanoparticles, gadolinium(III)-oxide ( $\text{Gd}_2\text{O}_3$ ) nanoparticles, europium(III)-oxide ( $\text{Eu}_2\text{O}_3$ ) nanoparticles, or terbium(III)-oxide ( $\text{Tb}_2\text{O}_3$ ) nanoparticles.

Embodiments synthesize Janus ceramic nanoparticles at mass quantity through one-pot reactions in a liquid-crystal lamellar phase. The liquid-crystal lamellar-phase medium may consist of surfactant, water, and organic compound(s) (for example, oil) to generate layered organic-water interfaces. The organic-water interfaces may be generated at intervals, for example, of about 10 nanometers (nm). Each organic-water interface (interfacial unit) can serve as an interfacial nanoreactor that provides interfacial reaction area for the reactions that form the Janus nanoparticles.

In implementations, chemical precursors are mixed in an organic phase solution or an aqueous phase solution. Then, the two solutions are mixed at controlled surfactant-water-organic ratios to form the liquid-crystal lamellar phase. Chemical reactions may occur upon mixing the precursors from the two phases. When the reactions are complete or near completion, Janus nanoparticles with one side hydrophilic and another side hydrophobic may be formed. Certain functional groups on the Janus nanoparticles, such as the amine functional group  $-\text{NH}_2$  and thiol functional group  $-\text{SH}$ , reserve opportunities for further functionalization that may be pursued depending on needs for various applications. Some embodiments are the direct formation of Janus nanoparticles without need for a post-synthesis surface functionalization. The direct formation may produce the Janus nanoparticles in quantities for industrial application.

A particular microstructure of interest is the organic lyotropic liquid crystal with long-range periodically structural order (the so-called lamellar ( $L\alpha$ ) phase) formed in part by a surfactant or mixtures of a surfactant and a co-surfactant in solvents. The lamellar phase consists of stacks of periodic molecular bilayer sheets separated by layers of solvents. By adding a solvents, such as water and an organic (for example, oil), the bilayer separation (d, lamellar periodicity) in the lyotropic lamellar phases can be swollen from one nanometer to tens of nanometer. A graphic illustration of the lamellar phase is displayed in FIG. 1.

Embodiments employ a lyotropic liquid-crystal phase as a template to produce Janus nanomaterials. The liquid-crystal phase medium may include surfactant, water, and an organic compound (oil) to generate layered organic-water interfaces with intervals in nanometers, such as approximately 10 nm. The repeating organic-water interfaces can amplify the interfacial area significantly by at least about a million times (for example, at about  $10^6$  order) in comparison to conventional bi-phase solution techniques. The present embodiments can perform interfacial reactions and facilitate production of the Janus nanosheets in quantities for industrial application.

Lamellar phase may refer generally to packing of polar-headed long chain nonpolar-tail molecules as sheets of bilayers separated by bulk polar liquid. As mentioned, a particular microstructure for certain embodiments is the organic lyotropic liquid-crystal with periodically structural order. This microstructure may be labeled as the lamellar ( $L\alpha$ ) phase or smectic A mesophase and formed in solvents (for example, water and organic compounds) with a surfactant or mixtures of surfactant and cosurfactant. The surfactant may be nonionic, anionic, cationic, or zwitterionic. The lamellar phase may consist of stacks of periodic molecular bilayer-sheets separated by layers of solvents. By adding solvents, such as water and an organic compound (for example, oil), the bilayer separation (for example, d-spacing or d, lamellar periodicity) in the lyotropic lamellar phases can be increased from a nanometer to tens of nanometer.

A liquid crystalline mesophase may be called lyotropic if formed by dissolving an amphiphilic mesogen in a solvent under specified conditions of concentration, temperature, and pressure. In contrast to thermotropic liquid crystals, lyotropics liquid crystals have an additional degree of freedom in the concentration that enables them to induce a variety of different phases. As the concentration of amphiphilic molecules is increased, several different type of lyotropic liquid crystal structures occur in solution, and the lamellar phase may be formed. This lamellar phase is denoted by the symbol  $L\alpha$  and can be considered the lyotropic equivalent of a smectic A mesophase. The lamellar phase may consist of amphiphilic molecules arranged in bilayer sheets separated by layers of water. A mesophase may be a state of matter intermediate between liquid and solid. Gelatin is a common example of a partially-ordered structure in a mesophase.

Surfactants may be compounds that reduce the surface tension or interfacial tension between two liquids, between a gas and a liquid, or between a liquid and a solid. With respect to the surfactants, the term "nonionic" may mean not ionic, not anionic, not cationic, and not zwitterionic. For a detergent, nonionic may mean that the detergent does not dissociate into ions in aqueous solution. A nonionic surfactant may have covalently bonded oxygen-containing hydrophilic groups, which are bonded to hydrophobic parent structures. The water-solubility of the oxygen groups may be the result of hydrogen bonding. Hydrogen bonding generally decreases with increasing temperature, and the water solubility of nonionic surfactants therefore decreases with increasing temperature.

FIG. 1 is a liquid-crystal lamellar phase **100** having a water phase (water layers **102**) and an organic phase (organic layers **104**). The lamellar phase **100** has alternating water layers **102** and organic layers **104**. The lamellar phase **100** includes a surfactant **106**. See FIG. 3 for examples of surfactants that may be utilized as the surfactant **106**. The multiple lyotropic liquid crystals of the lamellar phase **100** are the surfactant **106** molecules each having a hydrophilic head **108** and a hydrophobic tail **110** in the illustrated implementation. For clarity in the two water layers having the reference numeral **102**, only some of the hydrophilic heads **108** are depicted.

In some embodiments, the lamellar phase **100** may be prepared by dissolving a surfactant **106** (with or without a cosurfactant) into water and then adding an organic solvent (for example, oil or diesel) under agitation. The surfactant **106** can be a nonionic surfactant, a cationic surfactant, an anionic surfactant, or a zwitterionic surfactant. The surfactant **106** can be a mixture of a surfactant and a cosurfactant.

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The water layers **102** may be characterized as water phase layers **102**. The water layers **102** may be more generally an aqueous layer or hydrophilic layer. The water layers **102** may include ground water, treated water (for example, demineralized water), or formation water. The water layers **102** may include components other than water such as dissolved solids. The water layers **102** as an aqueous layer may include chemical components soluble in water. The water layers **102** have a thickness **112**.

The organic layers **104** may be characterized as organic phase layers **104**. The organic phase or organic layers **104** may be oil, such as mineral oil, refined oil, crude oil, fractions of crude oil, diesel, synthetic oil, or vegetable oil. The organic layers **104** may be, for example, heptane, octane, decane, dodecane, or diesel. The organic layers **104** are generally hydrophobic. The organic layers have a thickness **114**.

As mentioned for the illustrated embodiment, the surfactant **106** molecules each have a hydrophilic molecular head **108** and a hydrophobic molecular tail **110**. The lamellar phase **100** may typically be a planar structure with surfactant hydrophilic heads **108** contacting water layers **102** and surfactant hydrophobic tails **110** contacting the organic phase layers **104** in the illustrated implementation. In implementations, the lamellar phase **110** is generally not a micelle or emulsion.

In some instances, the organic layer **104** and water layer **102** may each be labeled as a lamellar layer. While the combination of the organic layer **104** and the water layer **102** (and with their interface) may instead be labeled as the lamellar layer, this combination is typically instead labeled as a lamellar unit **116** (d-spacing) as depicted. The lamellar unit **116** may include an organic layer **104**, a water layer **102**, the organic-water interface **118**, and the surfactant **106**. The surfactant hydrophilic heads **108** are on the water side of the interface **118**. The surfactant hydrophobic tails **110** are on the organic side of the interface **118**. The repeating lamellar unit **116** may have a thickness, for example, in a range of 5 nm to 100 nm. In certain embodiments, each lamellar unit **116** may be a nanoreactor.

Embodiments employ the lamellae as template media to produce Janus nanoparticles. The lamellar phase **100** can generate alternating interfaces of water and organic layers on the nanoscale range. The many lamellar units **116** may each be utilized respectively as a nanoreactor for chemical reactions at the water-organic interfaces **118**. The chemical reaction can be controllably-run occurring either at the water phase side or the organic phase side of the interface **118**, or both sides of the interface **118**. Employing these nanoreactors may provide to functionalize sides (surfaces) of the nanoparticles, producing Janus nanoparticles with one side hydrophilic and the other side hydrophobic. These nanoreactors may also be utilized to directly form the Janus nanoparticles via reaction of chemical precursors at the interface **118** without subjecting the surface of the nanoparticles to surface treatment, surface modification, or surface functionalization.

The lamellar phase **100** gives many water-organic interfaces **118** with each lamellar unit **116** being a respective lamellar nanoreactor. The lamellar phase **100** may have up to a million lamellar units **116** or more depending on the amount of surfactant and solution capacity. In the lamellar phase **100**, most or all of the chemical reagents added to the lamellar phase **100** may be confined in thin layers of solvents (organic layers **104** and water layers **102**) near the interfaces **118**. Therefore, chemical reactions may be more effective than in a bi-phase system having a single interface. In

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implementations, the chemical reactions in the lamellar phase **100** incorporate greater than 90 weight percent of the chemical reagents (chemical precursors) added to the lamellar phase **100**.

The lamellar phase **100** may provide for formation of Janus nanoparticles at the organic-water interfaces **118** in the lamellar phase **100**. As discussed, the alternating organic layers **104** and water layers **102** form a repeating organic-water interface **118**. These repeating organic-water interfaces may provide for repeating nanoreactors. Each lamellar unit **116** may include a single organic layer **104**, a single water layer **102**, the associated single organic-water interface **118**, and the associated surfactant **106** molecules having the hydrophilic heads **108** and hydrophobic tails **110**. The lamellar phase **100** may have at least one million lamellar units **116**. Thus, there may be at least one million nanoreactors in the lamellar phase **100**. In one implementation, the lamellar phase **100** has at least 100,000 lamellar units **116**, which may give a corresponding at least 100,000 nanoreactors.

For the lamellar phase **100**, the lamellar periodicity (d) may vary from several nanometers to several tens of nanometers by changing thickness of the organic (for example, oil) layers **104** or the water layers **102**, or both. This lamellar periodicity may be measured, for example, by small-angle x-ray scattering (SAXS) measurement or small-angle neutron scattering (SANS) measurement. The thickness of the water layers **102** in the lamellar phase **100** may be, for example, in the range of 5 nm to 10 nm. The thickness of the water layers **102** may be less than 50 nm, or less than 25 nm. The thickness of the organic layers **104** in the lamellar phase **100** may be, for example, in the range of 2 nm to 15 nm. The thickness of the organic layers **104** may be less than 50 nm, less than 30 nm, or less than 25 nm.

Embodiments may provide for the direct formation Janus nanoparticles in the lamellar template media. The lamellar phase **100** can generate organic-water interfaces **118** on the nanoscale range. Each lamellar unit **116** can be utilized as an interfacial nanoreactor for chemical reactions at the organic-water interface **118**. Chemical precursors (chemical reagents) dissolved in the water phase **102** and oil phase **104**, respectively, can meet at organic-water interface **118**. Therefore, chemical reactions occur at or near the organic-water interface **118** for nanoparticle formation. The hydrophobic components of the precursors prefer to stay at the oil phase side. The hydrophilic components of the precursors prefer to stay at the water phase side. Thus, the reactions from both hydrophilic and hydrophobic components at the interface may result in the direct formation Janus nanoparticles. A schematic illustration of Janus nanoparticle formation in the lamellar nanoreactors is shown in FIG. 2.

FIG. 2 is a sequence diagram **200** of a lamellar phase **100** in the formation of Janus nanoparticles **202**. Components mixed to form the lamellar phase **100** include water, an organic compound (for example, oil), and surfactant **106**. As discussed with respect to FIG. 1, the lamellar phase **100** includes alternating water layers **102** and organic layers **104**. As depicted in FIG. 2, chemical reagents **204** may be added to the lamellar phase **100**.

The chemical reagents **204** include chemical precursors **206** for the organic phase and chemical precursors **208** for the water phase. The chemical precursors **206** may be labeled as hydrophobic precursors. The chemical precursors **208** may be labeled as hydrophilic precursors. In some implementations, the hydrophobic precursors **206** may be organic-soluble (or oil-soluble) organometallic compounds and the hydrophilic precursors **208** may be water-soluble

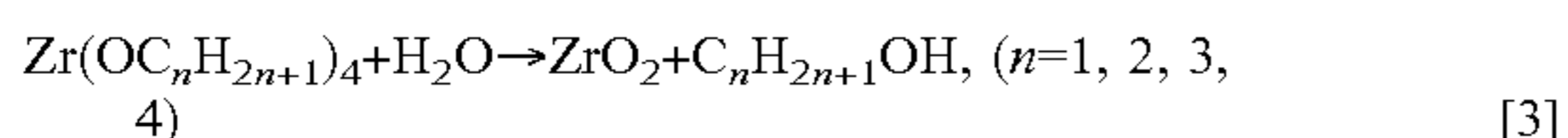
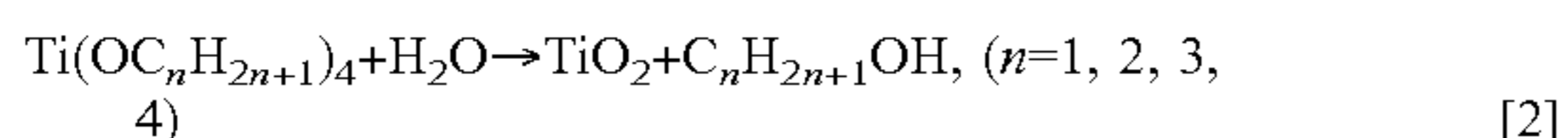
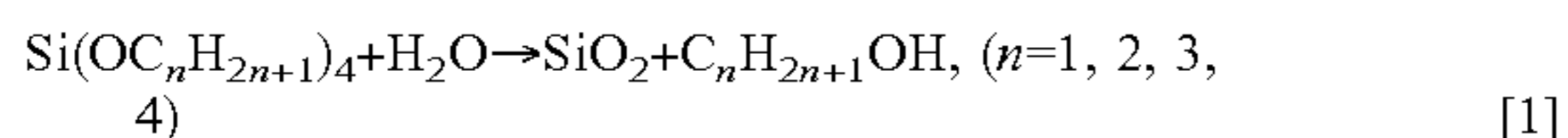
organometallic compounds. Solubility may be affected by saturation conditions of the lamellar phase **100**. In embodiments, at least 99 weight percent of the organic-soluble precursors **206** added may dissolve in the organic phase (layers **104**) of the lamellar phase **100**. At least 99 weight percent of the water-soluble precursors **208** may dissolve in the water phase (layers **102**) of the lamellar phase **100**.

In certain implementations, the chemical precursors **206** and **208** may be incorporated in an organic compound (for example, oil) and water, respectively, prior to addition to the lamellar phase **100**. As referenced by arrow **210**, the chemical precursors **206**, **208** may be added with the organic compound and water contemporaneous with the formation of the lamellar phase **100** or may be added after formation of lamellar phase **100**.

As referenced by arrow **212**, the chemical precursors **206**, **208** migrate toward the organic-water interfaces **118** where chemical reactions occur. A catalyst may be added to the lamellar phase **100** to accelerate the reactions. An example of a catalyst is ammonium hydroxide, tetramethylammonium hydroxide, hydrogen chloride, or acetic acid.

As referenced by arrow **214**, Janus nanoparticles **202** are formed at the organic-water interfaces **118** via the reaction of the chemical precursors **206** with the chemical precursors **208** at the organic-water interfaces **118**. In this implementation, the Janus nanoparticles **202** have a hydrophilic side **216** and a hydrophobic side **218**. The Janus nanoparticles **202** may straddle the interface **118**. The hydrophilic side **216** may reside in a water layer **102**. The hydrophobic side **218** may reside in an oil layer **104**. The Janus nanoparticles **202** directly formed may be Janus metal-oxide nanoparticles.

In the syntheses of Janus nanoparticles **212**, such as Janus SiO<sub>2</sub> nanoparticles, Janus TiO<sub>2</sub> nanoparticles, or Janus ZrO<sub>2</sub> nanoparticles, the hydrophilic precursors **208** may include water-soluble organometallic compounds. Examples include tetraalkyl orthosilicate (alkyl is methyl or ethyl), tetraalkyl orthotitanate (alkyl is ethyl, propyl, isopropyl, butyl, or tert-butyl), and tetrapropyl zirconate (alkyl is ethyl, propyl, isopropyl, butyl, or tert-butyl). These organometallic compounds can react with water to form metal oxide nanoparticles in aqueous phase (water layers **102**) with or without a catalyst, as depicted in equations [1], [2], and [3]:



In the syntheses of Janus metal-oxide nanoparticles (for example, Janus SiO<sub>2</sub> nanoparticles, Janus TiO<sub>2</sub> nanoparticles, or Janus ZrO<sub>2</sub> nanoparticles), examples of the hydrophobic precursors **206** may include oil-soluble organometallic compounds, such as n-alkyltriethoxysilane (alkyl is hexyl, octyl, decyl, dodecyl, tetradecyl, hexadecyl, or octadecyl), perfluoroalkyl-1H,1H,2H,2H-triethoxysilane (alkyl is hexyl, decyl, or octylphenyl), aminoalkyltriethoxysilane (alkyl=undecyl or octadecyl), and phenylalkyltrimethoxysilane, (alkyl is ethyl, propyl, or butyl). These organometallic compounds can reach an organic-water interface **118** and hydrolyze with water to form a thin layer on the metal-oxide nanoparticles while hydrophobic tails of these precursors **206** remain in the organic-phase side (in the organic layer **104**) at the organic-water interface **118**.

Synergetic effects of the reaction on the water side (water layer **102**) of the interface **118** with the reaction on the

organic side (organic layer **104**) of the interface **118** can result in formation of metal-oxide nanoparticles **212** with Janus structure. Exemplary photos of nanoparticles **212** as synthesized SiO<sub>2</sub> nanoparticles with different surface properties are shown in FIG. **4**. The asymmetric surface property of the Janus nanoparticles (in this instance, one side hydrophilic and another side hydrophobic) makes the Janus nanoparticles tend to aggregate at the water-organic interface **118**.

FIG. **3** gives exemplary surfactants **300** utilized to form the lamellar phase. The surfactants **300** to build the lamellar phase nanoreactors can be cationic, anionic, or nonionic. An example of a cationic surfactant is cetyltrimethylammonium bromide (CTAB) **302**. An example of an anionic surfactant is dioctyl sulfosuccinate sodium (AOT) **304**. An example of a nonionic surfactant is polyoxyethylene(4) lauryl ether (Brij® 30) **308**. In the growth of the Janus nanoparticles, a cationic surfactant can attract the forming nanoparticles (for example, ceramic nanoparticles) from the water phase to the water-organic interface via negative surface charges. However, nonionic and anionic surfactants may instead be employed.

## EXAMPLES

The Examples formed Janus SiO<sub>2</sub> nanoparticles. As discussed, the organic phase can be, for example, decane, dodecane, mineral oil, diesel, or crude oil. To facilitate characterization, decane was employed as the organic phase in the laboratory in the syntheses in the Examples. Diesel or crude oil as the organic layers may be utilized in the scale-up synthesis for commercial-scale application.

### Example 1

A first mixture was prepared by mixing (dissolving) the cationic surfactant CTAB and the hydrophilic precursor tetraethoxysilane (TEOS) in water. A second mixture was prepared by dissolving the hydrophobic precursor aminoundecyltriethoxysilane in decane. The first mixture and the second mixture were combined with a vortex mixer to form a generally uniform lamellar phase. Ammonium hydroxide as catalyst was added to the formed lamellar phase to accelerate the reaction. The chemical reactions in the lamellar phase occurred at room temperature for 12 hours without stirring. Upon the completion of the chemical reactions (at 12 hours of reaction), the lamellar phase was disabled by adding isopropanol. The formed Janus SiO<sub>2</sub> nanoparticles were collected by centrifuge and then washed by ethanol, acetone, and water in turn. Then, some of the Janus SiO<sub>2</sub> nanoparticles collected and washed were placed in a vial **406**, as depicted in FIG. **4**.

FIG. **4** is an image **400** of three vials **402**, **404**, and **406** each having different respective nanoparticles in hexane **410** and water **408**. The first vial **402** has hydrophilic pristine nanoparticles. As can be seen by the lack of clarity in the water phase **408** in the first vial **402**, the hydrophilic nanoparticles dispersed into the water phase. The second vial **404** has hydrophobic surface-functionalized nanoparticles. As can be seen by the lack of clarity in the hexane phase **410** in the second vial **404**, the hydrophobic nanoparticles dispersed into the hexane phase **410**. The third vial **406** has the Janus structure SiO<sub>2</sub> nanoparticles synthesized in Example 1. As can be seen by the lack of clarity at the water-hexane interface **412** in the third vial **406**, the Janus structure SiO<sub>2</sub> nanoparticles (formed in Example 1) migrated to the water-hexane interface **412**.

The described synthesis in Example 1 was repeated multiple times with the shape of the formed Janus structure SiO<sub>2</sub> nanoparticles controlled through the reactions. At less concentration of precursors in the water phase, the formed Janus nanoparticles had spherical shape. At greater concentration of the precursors in the water phase, the formed nanoparticles tended to be rod-like shape. Morphology of the synthesized SiO<sub>2</sub> nanoparticles formed in lamellar template media were characterized by scanning electron microscope (SEM).

FIG. 5 are SEM images 500 of Janus SiO<sub>2</sub> nanoparticles 502, 504, and 506 synthesized in Example 1 as having different respective shapes (controllable morphologies). The images 500 show that sizes of the Janus SiO<sub>2</sub> nanoparticles 502, 504, and 506 are at nanometer scale. The primary Janus SiO<sub>2</sub> nanoparticles 502 are generally spherical at a diameter of about 10-20 nm. The Janus SiO<sub>2</sub> nanoparticles 504 are also generally spherical but at a diameter of about 30-50 nm. Lastly, the Janus SiO<sub>2</sub> nanoparticles 506 are rod-like (cylindrical) with a diameter of about 30 nm and a length of about 200 nm.

FIG. 6 is a diagram of a reaction 600 giving formation of an isothiocyanate-amine complex 602. A dye molecule, such as Rhodamine B isothiocyanate (RhBITC) 604, having an isothiocyanate reactive group (—N=C=S) 606, is reactive towards a primary amine group (—NH<sub>2</sub>) 608 of a surface-modified nanoparticle 610 to form a stable fluorescent complex, such as the isothiocyanate-amine complex 602. The —NH<sub>2</sub> 608 may be a hydrophobic group or a hydrophilic group. In view of the reaction 600, RhBITC 604 was utilized to confirm that the SiO<sub>2</sub> nanoparticles formed in Example 1 had a Janus structure.

To confirm that SiO<sub>2</sub> nanoparticles formed in Example 1 had a Janus structure, the SiO<sub>2</sub> nanoparticles were mono-facial modified with hydrophobic undecylamino groups (—C<sub>11</sub>H<sub>23</sub>—NH<sub>2</sub>) to give a primary amine group (—NH<sub>2</sub>) 608 on the hydrophobic side of the SiO<sub>2</sub> nanoparticles. The “modification” to give the surface amine group 608 occurred directly in the formation of the SiO<sub>2</sub> nanoparticles via the reaction of the precursors. These mono-facial modified SiO<sub>2</sub> nanoparticles were dispersed in a water-hexane biphasic mixture. The mono-facial modified SiO<sub>2</sub> nanoparticles stayed as a thin layer at the water-hexane interface.

A dye solution having RhBITC 604 was injected into the water phase of the water-hexane bi-phase mixture. Then, after 2 hours, the dyed water phase was replaced with water not having the dye solution. The mono-facial modified SiO<sub>2</sub> nanoparticles at the water-hexane interface had no noticeable change in the mixture with or without the dye solution. This behavior (see also FIG. 7) indicates that there was no reaction 600 of the RhBITC 604 dye molecules (in the water phase) with the modified SiO<sub>2</sub> nanoparticles at the hexane-water interface. The hydrophobic undecylamino groups (having the —NH<sub>2</sub>) on the modified surface of the mono-facial modified SiO<sub>2</sub> nanoparticles did not reach the RhBITC 604 dye molecules (in the water phase) and therefore no reaction 600 occurred. The plain surface (not having the hydrophobic —NH<sub>2</sub> group) of the mono-facial modified SiO<sub>2</sub> nanoparticles was on the water side of the hexane-water interface.

#### Example 2

To further confirm that SiO<sub>2</sub> nanoparticles synthesized in present implementations have a Janus structure, additional SiO<sub>2</sub> nanoparticles were synthesized and the —NH<sub>2</sub> group 608 placed on the hydrophilic surface of the SiO<sub>2</sub> nanopar-

ticles. Thus, these SiO<sub>2</sub> nanoparticles synthesized in Example 2 were surface modified (directly in formation) to have a hydrophilic —NH<sub>2</sub> group.

The SiO<sub>2</sub> nanoparticles in Example 2 were synthesized as follows. A first mixture was prepared by mixing (dissolving) the nonionic surfactant CTAB and the hydrophilic precursors TEOS and (3-aminopropyl)triethoxysilane (APTES) in water. A second mixture was prepared by dissolving the hydrophobic precursor octadecyltriethoxysilane in dodecane. The first mixture and the second mixture were combined with a vortex mixer to form a generally uniform lamellar phase. Ammonium hydroxide as catalyst was added to the mixture to accelerate the reaction that formed the Janus SiO<sub>2</sub> nanoparticles at the interface of the dodecane and the water.

In their formation via the reaction of the precursors, these SiO<sub>2</sub> nanoparticles in Example 2 were altered to be mono-facial modified SiO<sub>2</sub> nanoparticles. A hydrophilic compound having the propylamine (—C<sub>3</sub>H<sub>7</sub>—NH<sub>2</sub>) group was added to modify (functionalize) the hydrophilic surface of the nanoparticles with the —C<sub>3</sub>H<sub>7</sub>—NH<sub>2</sub> group so that the nanoparticles would have a —NH<sub>2</sub> group 608 that is hydrophilic (on the hydrophilic side of the nanoparticles). A hydrophobic compound having the octadecyl (—C<sub>17</sub>H<sub>35</sub>—CH<sub>3</sub>) group was added to modify (functionalize) the hydrophobic surface of the nanoparticles with the C<sub>17</sub>H<sub>35</sub>—CH<sub>3</sub> group. These Janus SiO<sub>2</sub> nanoparticles in Example 2 as surface modified were collected and dispersed in a water-hexane biphasic mixture. The mono-facial modified SiO<sub>2</sub> nanoparticles stayed as a thin layer at the water-hexane interface.

A dye solution having RhBITC 604 was injected into the water phase of the water-hexane bi-phase mixture. The hexane-water interface turned fluorescence (see also FIG. 7) during the injecting of the RhBITC 604 solution into the water phase, implying that the RhBITC molecules reacted (bonded) with the hydrophilic —NH<sub>2</sub> groups of the Janus SiO<sub>2</sub> nanoparticles from the water phase side. The hydrophilic (—C<sub>3</sub>H<sub>7</sub>—NH<sub>2</sub>) group having the —NH<sub>2</sub> on the modified surface of the mono-facial modified SiO<sub>2</sub> nanoparticles interacted with the RhBITC 604 dye molecules (in the water phase) and therefore the reaction 600 occurred. The plain surface (hydrophobic and not having the —NH<sub>2</sub> group) of the mono-facial modified SiO<sub>2</sub> nanoparticles was on the hexane side of the hexane-water interface.

The aforementioned use of the RhBITC 604 solution in Example 1 and Example 2 revealed the asymmetric properties of the SiO<sub>2</sub> nanoparticles and confirmed their Janus structure. This use of the RhBITC 604 solution in Example 1 and Example 2 was repeated in a chloroform-water bi-phase mixture having dye RhBITC 604 in the water phase, as indicated in FIG. 7.

FIG. 7 is an image 700 of two vials 702, 704 each having a bi-phase mixture of chloroform 706 and water 708. The water 708 phase includes the dye RhBITC 604 (see FIG. 6). The bi-phase mixture in the first vial 702 has the Janus SiO<sub>2</sub> nanoparticles of Example 1 with the hydrophobic —NH<sub>2</sub> group. The bi-phase mixture in the second vial 704 has the Janus SiO<sub>2</sub> nanoparticles of Example 2 with the hydrophilic —NH<sub>2</sub> group. The Janus SiO<sub>2</sub> nanoparticles in each vial 702 and 704 generally reside at the water-chloroform interface 710.

As can be seen in FIG. 7, the dye RhBITC 604 in the first vial 702 is dispersed through the water 708 phase, indicating that the RhBITC 604 did not react (bond) with the nanoparticles in the first vial 702. The reaction 600 (see FIG. 6) did not occur in the first vial 702. This was expected because the

—NH<sub>2</sub> group was on the hydrophobic side of the nanoparticles facing the chloroform **706** phase in the first vial **702**. In contrast, the dye RhBITC **604** in the second vial **704** is seen as not dispersed through the water **708** phase but concentrated onto the Janus nanoparticles at the interface, indicating that the RhBITC **604** reacted (bonded) (reaction **600**) with the nanoparticles at the water-chloroform interface **710** in the second vial **704**. This was expected because the —NH<sub>2</sub> group was on the hydrophilic side of the nanoparticles facing the water **708** phase in the second vial **704**. The image **700** of the two vials **702**, **704** having the described respective mixtures confirm that the SiO<sub>2</sub> nanoparticles are asymmetric and have a Janus structure.

Instead of the dye molecules, different functional molecules can be grafted to one side of the Janus nanoparticles, as noted in FIG. **8**. The Janus nanoparticles may be Janus metal-oxide nanoparticles. The attachment of functional molecules to the Janus nanoparticle can extend applicability of the Janus nanoparticles to multiple industries, such as applications in sensing and biomedical materials.

FIG. **8** is a diagram **800** depicting exemplary reactions that may further functionalize the Janus nanoparticles **802**. In these implementations, the Janus nanoparticles **802** have a functional group, such as an amine —NH<sub>2</sub> group **804**, on one side for the functionalization reaction to occur. The —NH<sub>2</sub> group **804** may be on the hydrophobic side of the Janus nanoparticles **802** or on the hydrophilic side of the Janus nanoparticles **802**.

The further functionalization of the Janus nanoparticles **802** with different functional groups may provide for applicability of the Janus nanoparticles in a range of applications. The conjugation schemes give attachment of molecules (R) onto the amine-coated surface of the Janus nanoparticles **802**. The molecules may include biomolecules. The molecules (R) are attached to the nanoparticle surface on the side of the Janus nanoparticle **802** having the amine coating (having the —NH<sub>2</sub> group **804**). The molecules (R) each have and present a target site (a functional group) for interaction or reaction with the amine group **804** on the nanoparticle **802** to attach or graft the molecule (R) to the surface of the nanoparticle **802**. The target sites (functional groups) may include, for example, an amine **806**, a thiol **808**, a hydroxyl **810**, a carboxyl **812**, and an isothiocyanate **814**. Exemplary reagents in the reactions may include 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide hydrochloride (EDC), succinimidyl ester (NHS), succinimidylidodoacetate (SIA), N-succinimidyl 3-(2-pyridyldithio) propionate (SPDP), succinic anhydride, and thionyl chloride (SOCl<sub>2</sub>).

Langmuir-Blodgett technology was employed to examine interfacial properties of the Janus nanoparticles. When injecting a suspension of Janus SiO<sub>2</sub> nanoparticles suspension into a water-hexane mixture, a nanoparticle monolayer forms at the water-hexane interface. In applying lateral pressure to compress the nanoparticle monolayer, the surface pressure ( $\pi$ ) can reach 20 millinewton per meter (mN/m) or greater, further confirming the formation of a stable Janus structure.

The Janus nanoparticles have an effect on a water-organic interface. When injecting a suspension Janus nanoparticles into a water-organic mixture, the interfacial tension at the water-oil interface can be altered, as indicated in FIG. **9**.

FIG. **9** is an image **900** of three vials **902**, **904**, and **906**. The first vial **902** has water **908** and crude oil **910** to depict a plain interface **912** of the water **908** and crude oil **910**. No nanoparticles are included in the bi-phase mixture in the first vial **902**. The second vial **904** has water **914** and a combination of crude oil and chloroform as the organic phase **916**.

The bi-phase mixture in the second vial **904** includes Janus SiO<sub>2</sub> nanoparticles. The water-organic interface **918** in the second vial **904** is more flat than the interface **912** in the first vial **902**. The third vial **906** has water **920** and crude oil **922**. The bi-phase mixture in the third vial **906** includes Janus SiO<sub>2</sub> nanoparticles. The water-crude oil interface **924** in the third vial **906** is more flat than the water-crude oil interface **912** in the first vial **902**.

Embodiments provide for a one-pot synthesis of Janus ceramic (metal oxide) nanoparticles at greater than one kilogram per day. The Janus metal-oxide nanoparticles can alter interfacial properties of water-organic (oil) interfaces. Implementations may provide for direct formation of Janus nanoparticles through a one-pot reaction at interfacial nanoreactors. The interfacial nanoreactors may be composed of lyotropic liquid-crystal lamellar phase for the synthesis. The chemical reaction routes for the ceramic nanoparticles formation may be through hydrolysis of organometallic compounds at the interfacial nanoreactors. Janus nanoparticles production in lyotropic liquid-crystal lamellar phase was demonstrated in the Examples. Structural and property characterizations on the synthesized Janus nanomaterials were performed.

Implementations include equipment, operating conditions, and production rate in commercial systems to produce these Janus metal-oxide nanoparticles at industrial scale. In embodiments, the chemical reactions occur at ambient temperature and atmosphere pressure. The quantity of product in one-batch reaction may depend on the volume of the reaction vessel. The scale can be amplified from lab scale in milliliters to intermediate scale at liters and industrial scale at tens or hundreds of liters.

Embodiments provide synthesizing millions of Janus nanoparticles in a single batch in a vessel. The technique may be at industrial scale and for a range of applications of Janus nanoparticles in various industries. The Janus nanoparticles can alter interfacial properties of water-oil interfaces. The Janus nanoparticles may have application in a nanoagent solution for enhanced oil recovery (EOR) in oil and gas recovery applications.

Embodiments employ lyotropic liquid crystal phase as a template for Janus nanoparticle synthesis and with efficiency of interfacial reactions by a million times (at 10<sup>6</sup> order) compared with bi-phase reactions. Such promotes and facilitate bulk Janus nanosheets production feasible for industrial applications. The nanoreactor utilizes a lyotropic liquid crystal phase as template media for chemical reactions at water-oil interfaces. A chemical reaction route is implemented to fabricate Janus nanoparticles in mass or bulk quantities and also to functionalize the Janus nanoparticles. The structure and properties of the synthesized Janus nanoparticles may be evaluated. Functionalized Janus nanoparticles may be characterized. Microfluidic imaging of the Janus nanoparticles may be performed for EOR applications.

FIG. **10** is a method **1000** of synthesizing Janus material, such as Janus nanomaterial or Janus nanoparticles. The Janus material is synthesized in a lamellar media or lamellar phase. The synthesis of the Janus nanoparticles may be performed in a vessel, such as in a batch operation. Semi-batch or continuous operations may be employed.

At block **1002**, the method includes forming or providing a lamellar phase having water layers and organic layers. The lamellar phase may typically include a surfactant. A surfactant may organize at a liquid-liquid boundary which can lead to an organized interface and liquid-crystal phases. The surfactant may be a nonionic surfactant, a cationic surfac-

tant, an anionic surfactant, or a zwitterionic surfactant. In certain implementations, hydrophilic molecular heads of the surfactant contact the water layers and hydrophobic molecular tails of the surfactant contact the organic layers. For a surfactant molecule, the molecule may have a hydrophilic head contacting a water layer and a hydrophobic tail contacting an organic layer adjacent the water layer. The lamellar phase may additionally include a cosurfactant. A cosurfactant may be a chemical utilized with a surfactant to improve performance or enhance effectiveness of the surfactant. An example of a cosurfactant is an alcohol having at least 5 carbons.

The water layers may be characterized as water phase layers. Likewise, the organic layers may be characterized as organic phase layers. The organic phase may be an oil phase and thus the organic layers may be oil layers. The lamellar phase may include at least 100,000 water layers and at least 100,000 organic layers. The water layers may alternate with the organic layers. The water layers may each have a thickness of up to about 50 nm. The organic layers may each have a thickness of up to about 50 nm. In certain embodiments, the lamellar phase is placed or formed in the same vessel in which the synthesis (for example, batch synthesis) of the Janus nanoparticles occurs.

The lamellar phase also includes chemical precursors (chemical reagents). The chemical precursors may be added to or incorporated in the lamellar phase after the lamellar phase is formed. The chemical precursors may be incorporated in the lamellar phase as the lamellar phase is being formed. The chemical precursors may be added contemporaneous with the formation of the lamellar phase. The chemical precursors may be included in the water and organic compound prior to mixing of the water, organic compound, and surfactant that forms the lamellar phase. Examples of the chemical precursors include organometallic compounds.

At block **1004**, the method includes reacting the chemical precursors in the lamellar phase to form Janus nanoparticles at interfaces of the water layers with the organic layers. The method may include allowing the chemical precursors to migrate to the interfaces where reactions between chemical reagents occur. The forming (block **1002**) of the lamellar phase and reacting (block **1004**) of the chemical precursors may to form the Janus nanoparticles may be performed in the same vessel as a one-pot synthesis.

The Janus nanoparticles may be Janus ceramic nanoparticles or Janus metal-oxide nanoparticles. The Janus metal-oxide nanoparticles may be, for example, Janus SiO<sub>2</sub> nanoparticles, Janus TiO<sub>2</sub> nanoparticles, or Janus ZrO<sub>2</sub> nanoparticles. The Janus nanoparticles may have a hydrophobic surface and a hydrophilic surface. At the interfaces, a hydrophilic surface or hydrophilic portion of the Janus nanoparticles may reside in the water layers. A hydrophobic surface or hydrophobic portion of the Janus nanoparticles may reside in the respective organic layer adjacent the water layer.

The reacting of chemical precursors may include reacting at the interfaces a water-soluble chemical precursor (reagent) incorporated in the water layers with an organic-soluble chemical precursor (reagent) incorporated in the organic layers. The water-soluble chemical precursor may be, for example, a water-soluble organometallic compound. The organic-soluble precursor may be, for example, an oil-soluble organometallic compound.

The reacting of the chemical precursors may give a surface functional group, such as an alkyl group (—C<sub>n</sub>H<sub>2n+1</sub>CH<sub>3</sub> group), an amine group (—NH<sub>2</sub> group), a thiol group

(—SH group), a carboxyl group (—COOH group) or an isothiocyanate group (—NCS group) on a surface of the Janus nanoparticles. The chemical precursors may include the functional group that becomes the surface functional group. The water-soluble precursor may have a functional group that becomes a functional group on a hydrophilic portion of the Janus nanoparticles. If so, the water-soluble precursor reacting with the organic-soluble precursor gives the functional group as a surface functional group on a hydrophilic side of the Janus nanoparticles. The organic-soluble precursor may have a functional group that becomes a functional group on a hydrophobic portion of the Janus nanoparticles. In these implementations, the organic-soluble precursor reacting with the water-soluble precursor gives the functional group as a surface functional group on a hydrophobic side of the Janus nanoparticles.

The surface functional group (whether on a hydrophilic surface or hydrophobic surface, or both) of the Janus nanoparticles may be, for example, an amine (—NH<sub>2</sub>) group or thiol (—SH) group. A purpose of the surface functional group may be for further functionalization of the Janus nanoparticles, as discussed with respect to block **1008**.

At block **1006**, the method includes collecting the Janus nanoparticles from the lamellar phase. For instance, the method may include centrifuging the lamellar phase to separate the Janus nanoparticles from the lamellar phase for collection. The collected Janus nanoparticles may be washed with water or a hydrocarbon.

At block **1008**, the method includes reacting a molecule functional group of molecules with a surface functional group (for example, discussed in block **1004**) in presence of a reagent to attach the molecule to a surface of the Janus nanoparticles. In other words, the method may include reacting a compound functional group of a compound with the surface functional group to attach the compound to the hydrophilic side of the Janus nanoparticles or to the hydrophobic side of the Janus nanoparticles, or both sides. The compound or molecule functional group may be an amine group, a thiol group, a hydroxyl group, a carboxyl group, or an isothiocyanate group, or any combinations thereof. The reagent may include EDC, NHS, SIA, SPDP, succinic anhydride, or SOCl<sub>2</sub>, or any combinations thereof.

An embodiment is a nanoreactor that includes an interface between a water layer of a lamellar phase and an organic layer (for example, oil layer) of the lamellar phase. The lamellar phase includes a surfactant. The nanoreactor may be characterized as including the water layer and the organic layer. The nanoreactor may be characterized as the lamellar unit including the water layer, organic layer, and water-organic interface. The nanoreactor includes Janus metal-oxide nanoparticles as product formed at the interface from chemical precursors in the lamellar phase. The chemical precursors include a water-soluble organometallic compound and an organic-soluble organometallic compound. The lamellar phase may include at least 100,000 lamellar units and thus may include at least 100,000 nanoreactors.

Another embodiment is a lamellar phase having multiple nanoreactors, each nanoreactor being an interface between a water layer (of the lamellar phase) and an organic layer (of the lamellar phase). In operation, the nanoreactors react a first chemical reagent that is water soluble with a second chemical reagent that is organic soluble to form Janus metal-oxide nanoparticles at the interface. A hydrophilic surface of the Janus nanoparticles resides in the water layer. A hydrophobic surface of the Janus nanoparticles resides in the organic layer. The lamellar phase has multiple water layers and multiple organic layers. A thickness of each water



layer in the lamellar phase may be less than 25 nm. A thickness of each organic layer in the lamellar phase may be less than 30 nm.

Yet another embodiment is a method of synthesizing Janus nanoparticles, such as Janus metal-oxide nanoparticles. The Janus nanoparticles may have a hydrophobic surface and a hydrophilic surface. The method includes forming a lamellar phase having water layers, organic layers, and a surfactant. A molecule of the surfactant may have a hydrophilic head contacting a water layer and a hydrophobic tail contacting an organic layer adjacent the water layer. The method includes reacting chemical precursors (for example, organometallic compounds) in the lamellar phase to form Janus nanoparticles at interfaces of the water layers with the organic layers. The reacting of the chemical precursors may involve reacting at the interfaces a first chemical precursor (for example, a water-soluble organometallic compound) incorporated in the water layers with a second chemical precursor (for example, an oil-soluble organometallic compound) incorporated in the organic layers. The forming of the lamellar phase and the reacting of the chemical precursors to form the Janus nanoparticles may be a one-pot synthesis.

The reacting of the chemical precursors may give a surface functional group (for example, an amine group or a thiol group) on a surface of at least one Janus nanoparticle of the Janus nanoparticles. The surface functional group may be, for example, an alkyl  $-C_nH_{2n+1}CH_3$  group, an amine  $-NH_2$  group, a thiol  $-SH$  group, a carboxyl  $-COOH$  group, or an isothiocyanate  $-NCS$  group. The method may include reacting a molecule functional group of a molecule with the surface functional group in presence of a reagent to attach the molecule to the surface of the Janus nanoparticle. The molecule functional group may include, for example, an amine group, a thiol group, a hydroxyl group, a carboxyl group, or an isothiocyanate group, or any combinations thereof. The reagent may include, for example, 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide hydrochloride (EDC), succinimidyl ester (NHS), succinimidylidodoacetate (SIA), N-succinimidyl 3-(2-pyridyldithio) propionate (SPDP), succinic anhydride, or thionyl chloride (SOCl<sub>2</sub>), or any combinations thereof.

Yet another embodiment is a method of synthesizing Janus nanoparticles, comprising forming a lamellar phase having a surfactant, water layers, and organic layers. The water layers may be at least 100,000 water layers each having a thickness less than 50 nm. The organic layers may be at least 100,000 organic layers each having a thickness less than 50 nm. The forming of the lamellar phase includes incorporating a first chemical reagent in the water layers and incorporating a second chemical reagent in the organic layers. The method includes allowing the first chemical reagent to react with the second chemical reagent at interfaces of the water layers with the organic layers to form Janus nanoparticles (for example, Janus ceramic nanoparticles) at the interfaces. A hydrophilic surface of a Janus nanoparticle of the Janus nanoparticles may reside in a water layer. A hydrophobic surface of the Janus nanoparticle may reside in an organic layer adjacent the water layer. The forming of the lamellar phase and allowing the first chemical reagent to react with the second chemical reagent may occur in the same vessel.

Yet another embodiment is a method of synthesizing Janus nanomaterial. The method includes forming a lamellar phase having a surfactant, water layers, organic layers (for example, oil layers), and chemical precursors. The method includes allowing the chemical precursors to react to form

Janus nanoparticles at interfaces between the organic layers and the water layers. A hydrophobic portion of the Janus nanoparticles may reside in the organic layers. A hydrophilic portion of the Janus nanoparticles may reside in the water layers. For instance, a hydrophobic portion of a Janus nanoparticle of the Janus nanoparticles may reside in an organic layer of the organic layers. A hydrophilic portion of the Janus nanoparticle may reside in a water layer of the water layers and where the organic layer is adjacent the water layer. The Janus nanoparticles may be, for example, Janus silicon-dioxide (SiO<sub>2</sub>) nanoparticles, titanium-dioxide (TiO<sub>2</sub>) nanoparticles, or zirconium-dioxide (ZrO<sub>2</sub>) nanoparticles, or any combinations thereof.

The chemical precursors may include a first chemical precursor and a second chemical precursor. The first chemical precursor may be a water-soluble chemical reagent. The second chemical precursor may be an oil-soluble chemical reagent. The allowing of the chemical precursors to react may involve the first chemical precursor reacting with the second chemical precursor. In implementations, the first chemical precursor has a functional group and where the first chemical precursor reacting with the second chemical precursor gives the functional group as a surface functional group on a hydrophilic side of a Janus nanoparticle of the Janus nanoparticles. The method may include collecting the Janus nanoparticle from the lamellar phase and reacting a compound functional group of a compound with the surface functional group to attach the compound to the hydrophilic side of the Janus nanoparticle. In implementations, the second chemical precursor may have a functional group and where the first chemical precursor reacting with the second chemical precursor gives the functional group as a surface functional group on a hydrophobic side of a Janus nanoparticle of the Janus nanoparticles. The method may include collecting the Janus nanoparticle from the lamellar phase and reacting a compound functional group of a compound with the surface functional group to attach the compound to the hydrophobic side of the Janus nanoparticle.

Yet another embodiment is a lamellar phase including surfactant, water layers having a water-soluble organometallic compound, and oil layers having an oil-soluble organometallic compound. The lamellar phase has Janus metal-oxide nanoparticles formed from reaction of the water-soluble organometallic compound with the oil-soluble organometallic compound at the interfaces of the water layers with the oil layers. The Janus metal-oxide nanoparticles are disposed at the interfaces. In implementations, a hydrophilic surface of a Janus metal-oxide nanoparticle of the Janus metal-oxide nanoparticles resides in a water layer and a hydrophobic surface of the Janus metal-oxide nanoparticle resides in an oil layer adjacent the water layer. The water layers may include at least 100,000 water layers. The oil layers may include at least 100,000 oil layers. The water layers may each have a thickness up to about 50 nm. The oil layers may each have a thickness up to about 50 nm.

A number of implementations have been described. Nevertheless, it will be understood that various modifications may be made without departing from the spirit and scope of the disclosure.

What is claimed is:

1. A method of synthesizing Janus nanoparticles, comprising:
  - forming a lamellar phase comprising water layers, organic layers, and a surfactant; and
  - reacting chemical precursors in the lamellar phase to form Janus nanoparticles at interfaces of the water layers with the organic layers, wherein reacting the chemical

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precursors gives a surface functional group on a surface of a Janus nanoparticle of the Janus nanoparticles; and reacting a molecule functional group of a molecule with the surface functional group in presence of a reagent to attach the molecule to the surface of the Janus nanoparticle, wherein the reagent comprises 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide hydrochloride (EDC), succinimidyl ester (NHS), succinimidylacetate (SIA), N-succinimidyl 3-(2-pyridyldithio) propionate (SPDP), succinic anhydride, or thionyl chloride (SOCl<sub>2</sub>), or any combinations thereof.

2. The method of claim 1, wherein a molecule of the surfactant comprises a hydrophilic head contacting a water layer and a hydrophobic tail contacting an organic layer adjacent the water layer, and wherein the chemical precursors comprise organometallic compounds.

3. The method of claim 1, wherein reacting chemical precursors comprises reacting at the interfaces a first chemical precursor incorporated in the water layers with a second chemical precursor incorporated in the organic layers.

4. The method of claim 3, wherein the first chemical precursor comprises a water-soluble organometallic com-

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pound, wherein the second chemical precursor comprises an oil-soluble organometallic compound, and wherein the Janus nanoparticles comprise Janus metal-oxide nanoparticles.

5. The method of claim 1, wherein the Janus nanoparticles comprise a hydrophobic surface and a hydrophilic surface, and wherein forming the lamellar phase and reacting the chemical precursors to form the Janus nanoparticles is a one-pot synthesis.

6. The method of claim 1, wherein the surface functional group comprises an amine group or a thiol group.

7. The method of claim 1, wherein the surface functional group comprises an alkyl —C<sub>n</sub>H<sub>2n+1</sub>CH<sub>3</sub> group, an amine —NH<sub>2</sub> group, a thiol —SH group, a carboxyl —COOH group, or an isothiocyanate —NCS group.

8. The method of claim 1, wherein the molecule functional group comprises an amine group, a thiol group, a hydroxyl group, a carboxyl group, or an isothiocyanate group, or any combinations thereof.

\* \* \* \* \*

UNITED STATES PATENT AND TRADEMARK OFFICE  
**CERTIFICATE OF CORRECTION**

PATENT NO. : 11,572,282 B2  
APPLICATION NO. : 16/653367  
DATED : February 7, 2023  
INVENTOR(S) : Wei Wang

Page 1 of 1

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

On the Title Page

Column 2, Line 12, please replace "Febmaty" with -- February --.

Column 2, Line 14, please replace "lemellar" with -- lamellar --.

Column 2, Line 21, please replace "inorganicnanoparticles" with -- inorganic nanoparticles --.

In the Claims

In Column 18, Line 16, Claim 7, please replace "isothiolcyanate" with -- isothiocyanate --.

Signed and Sealed this  
Thirtieth Day of May, 2023  
*Katherine Kelly Vidal*

Katherine Kelly Vidal  
*Director of the United States Patent and Trademark Office*